

Fig. 1

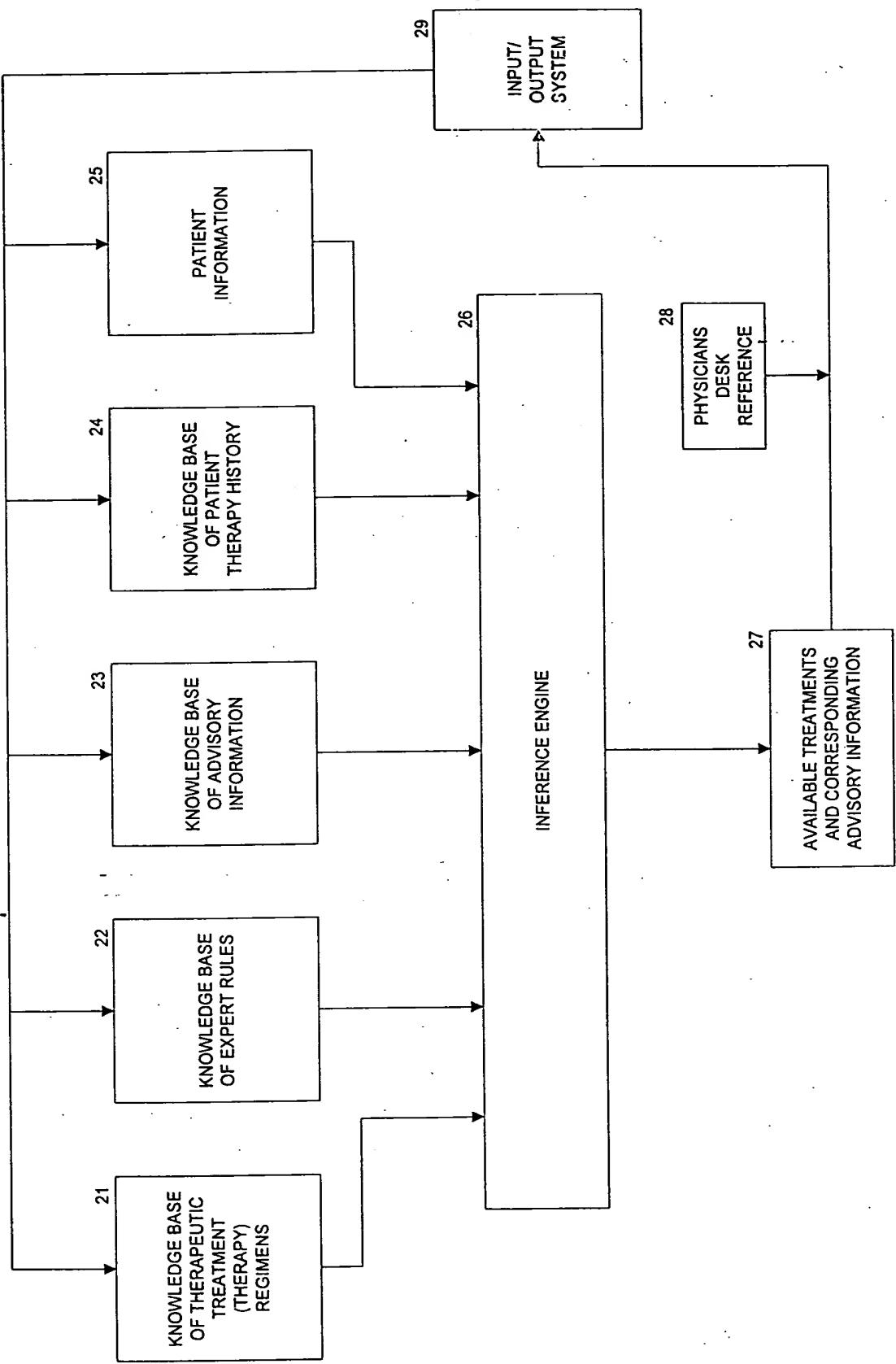
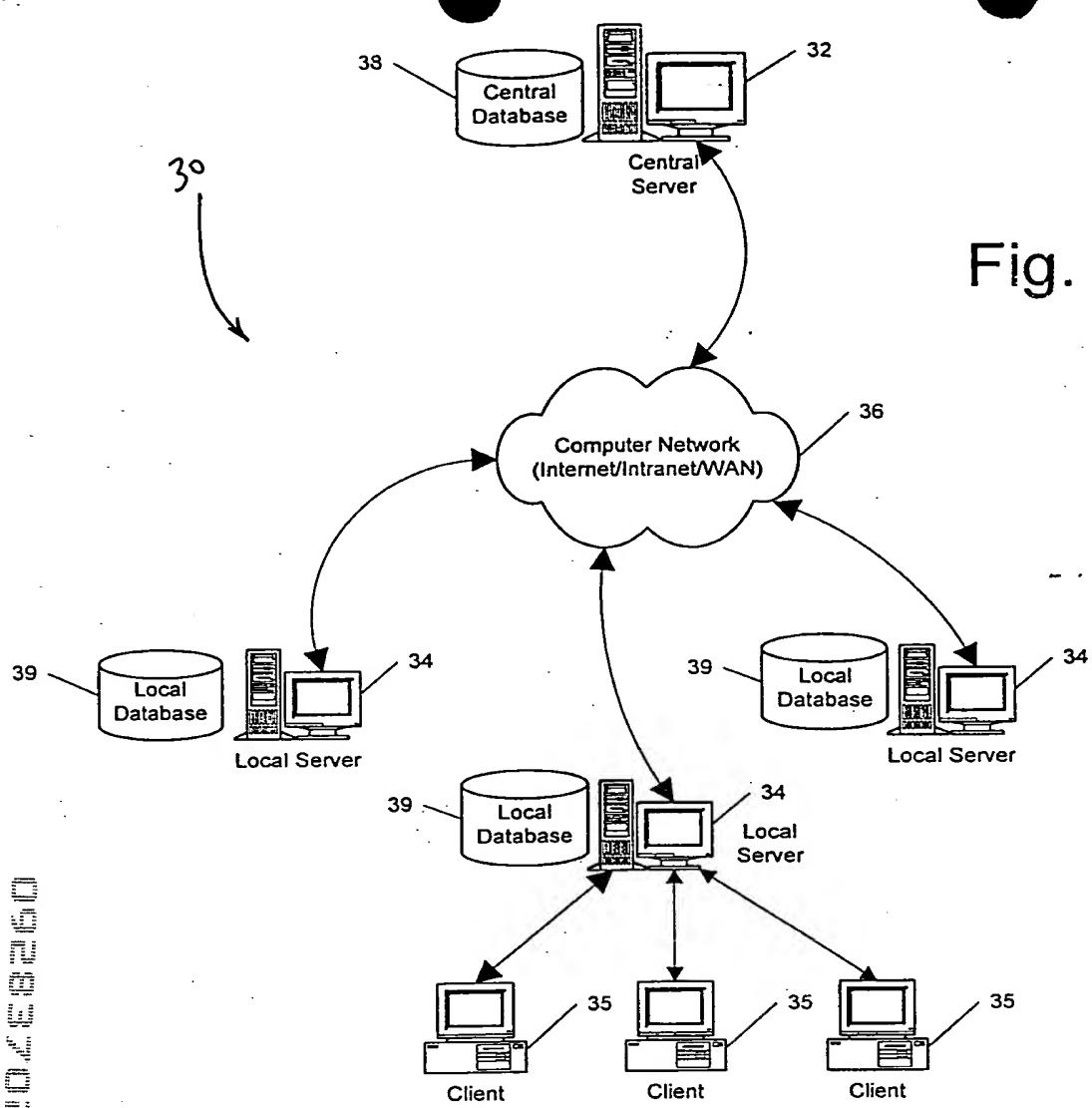


Fig. 2

Fig. 3



卷之三

F/16. 4

61

F16.5

61

63

62

64

63

64

F16.7
RECORDS

Icon	Meaning
◎	Indicates that there were no critical alerts for the therapy, however, general warnings and advisories should be read in the Therapy Details box.
●	Indicates that there were no critical alerts for the therapy, however, general warnings and advisories should be read in the Therapy Details box. The book indicates that therapy has been studied and a reference is available to review.
△	Indicates a yellow alert. There is important information about this therapy that must be reviewed.
■	Indicates a yellow alert. There is important information about this therapy that must be reviewed. The book indicates that therapy has been studied and a reference is available to review.
!	Indicates a red alert, which means critical and possible life-threatening situation may exist or may be created with this therapy. Information in the Therapy Details section must be read for this therapy to be considered.
●	Indicates a red alert, which means critical and possible life-threatening situation may exist or may be created with this therapy. Information in the Therapy Details section must be read for this therapy to be considered. The book indicates that therapy has been studied and a reference is available to review.
✗	Indicates the therapy is not recommended.

卷之三

72a

736 78

73

73d

73e

73f

734

四

1003

- Rituximab 300mg q12h (2 pills/day, \$9,56/day)
- Vidarabine 125mg q12h (4 pills/day, \$4,22/day)
- Lamivudine 400mg q12h; taken within 2 hours after a full meal (4 pills/day, \$8,47/day)
- Nevirapine 200mg q12h (10 pills/day, \$1,43/day)

(\ddagger indicates adjusted dosage)

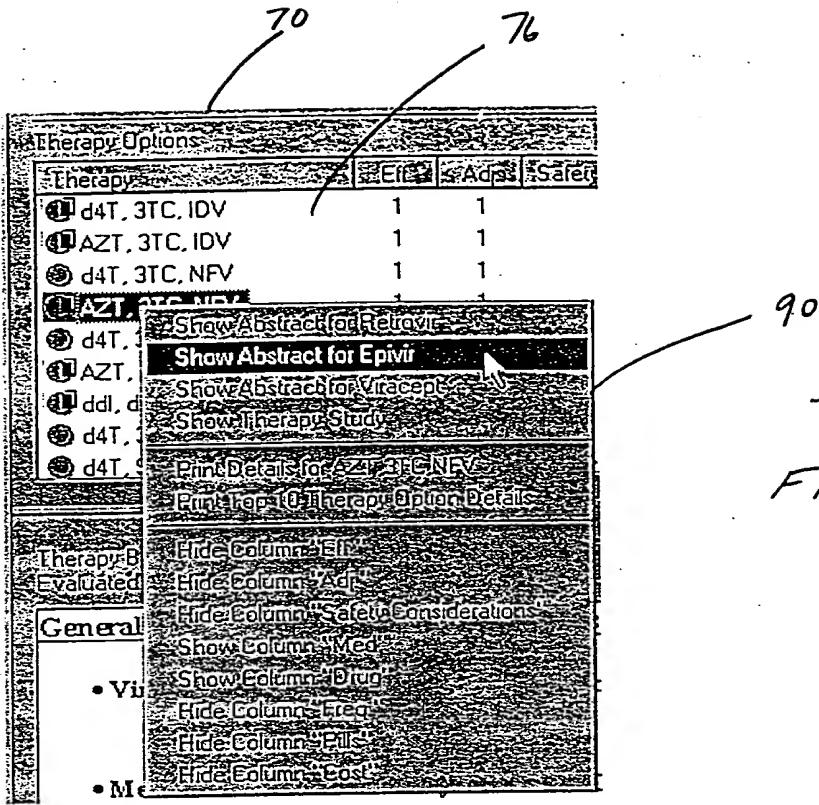
24

દ્વારા લખાયા શાખા હતી કે સુધીએ જોકાંના ગુણીયતાની વિષયે પ્રાચીન રીતે

Dosage Notice: This therapy contains both saquinavir and ritonavir. When ritonavir and saquinavir are used together the dosage of each drug is reduced by 1/2. The dosage

SAQ: The following Warnings and Advisories apply to Intrinsic (solid) materials.

- **Drug Interactions:** Compounds that are substrates of CYP2A6 (e.g., calcium channel blockers, clindamycin, dapsone, quinidine, triiodothyronine) may have elevated plasma concentrations when coadministered with lurasidone (quinuvin/SQV); therefore, patient should be monitored for toxicities associated with such drugs when taking lurasidone (quinuvin/SQV). *ComCat® Comment 2*



TPMS

/ Sea / 600 / 700

Ward/Hemo/Chu | Therapy/Evaluation

General		<input checked="" type="checkbox"/> F1	<input checked="" type="checkbox"/> F2	<input checked="" type="checkbox"/> F3	<input checked="" type="checkbox"/> F4	<input checked="" type="checkbox"/> F5
Patient ID	Date of Birth	TPMS Number	Specimen Date	Weight (kg)	Date	Value
demo1	1/1/1960	1000	3/1/1999	55.00	3/3/1999	55.00
Shoemaker	Gender	Male	Specimen	Solid/Dissolve	Specimen Date	Yes
CD4 and Viral Load						
CD4 (cells/mm ³)	Specimen Date	Value	Specimen Date	Prev Value	AIDS Defining Event	
100	3/1/1999	320	1/1/1999	140		
Current Viral Load	Specimen Date	Value	Specimen Date	Prev Value		
10000	3/1/1999	120000	1/1/1999	100000		
Previous Viral Load	Specimen Date	Value	Specimen Date	Prev Value		
500	1/1/1999	500	1/1/1999	500		
HIV Genotype						
Phenotype	Specimen Date	Value	Specimen Date	Prev Value	Last Drug Start Date	
Altered/Hyper	Specimen Date	Value	Specimen Date	Prev Value	1/1/1999	
Intolerance	Specimen Date	Value	Specimen Date	Prev Value		
Hemoglobin						
Specimen Date	Value (g/l)	Date	Date	Date	Last Drug Start Date	
3/1/1999	12.00	3/1/1999	No	No		
Neutrophil						
Specimen Date	Value (x10 ⁹ /L)	Date	Date	Date	Last Drug Start Date	
3/1/1999	1500	3/1/1999	No	No		
Hepatic Function						
Specimen Date	AST (SGOT) (U/L)	Date	Date	Date	Last Drug Start Date	
3/1/1999	49	45	3/1/1999	No	2.00	3/1/1999

F2

F3

F4

F5

546

547

F1

F16.10A

TPMS 41 2

How To

TPMS Patient

Medical History / Chart / Therapy Evaluation

Eveready, Ernestine 49y | XZ | 364-DV

Therapy Selection (0 of 38)

Therapy	Eff.	Adj.	Safety Considerations	Freq.	Pills	Dosage
ddI, d4T, NVP	2	2	ddI Renal dos. Adj. d4T Renal dos. adj	q8h	15	\$30.38
△ ddI, d4T, RTV	4	4	ddI Renal dos. Adj. d4T Renal dos. adj	q12h	18	\$34.06
△ NVP, ABC, EFV	5	5	NVP Renal dos. Adj. EFV+Renal Dysf	q12h	9	\$44.32
△ DLV, ABC, EFV	5	5	EFV+Renal Dysf	q8h	13	\$43.21
△ NVP, ABC, EFV	5	5	EFV+Renal Dysf	q8h	16	\$54.40
△ NVP, NVP, EFV	5	5	NVP Renal dos. Adj. EFV+Renal Dysf	q8h	17	\$46.41

Second | Therapy | Freq. | Pill Count | Evaluation

Tramadol | XZ | 364-DV
Endured

Therapy	Eff.	Adj.	Safety Considerations	Freq.	Pills	Dosage
Zidovudine XZ 364-DV	4	4	Zidovudine ZDV	q12h	18	\$34.06
△ Zidovudine XZ 364-DV	5	5	Zidovudine ZDV	q12h	18	\$34.06
△ Zidovudine XZ 364-DV	5	5	Zidovudine ZDV	q12h	18	\$34.06
△ Zidovudine XZ 364-DV	5	5	Zidovudine ZDV	q12h	18	\$34.06

- AZT△: Medical Condition Alert: This patient has a history of anemia. Use Retrovir with caution due to risk of hematologic toxicity. More Info 171

FiltRankC, Commentary 171

73

Recommended Dosages

- Retrovir 300mg q12h (2 pills/day, \$9.56/day)
- **Epivir** 150mg q24h (1 pill/day, \$3.84/day)
- Crixivan 800mg q8h (6 pills/day, \$15.00/day)

(+ indicates adjusted dosage)

Warning - Resistance Notices

[Redacted]

- Resistance Advisory: Retrovir and Epivir ranked lower (+2) due to historical virological failure. More Info 364 FiltResP13 Commentary 364

• Dolutegravir: Dosing adjustment to 50mg daily due to resistance to dolutegravir. Last outcome date: March 10, 2011

TPMS 41 1

10W WEB SW SW

70

Medical History / Clinical Therapy Evaluation																					
General	<input checked="" type="checkbox"/> HIV <input type="checkbox"/> HCV <input type="checkbox"/> CMV <input type="checkbox"/> EBV <input type="checkbox"/> Toxoplasma Patient ID: demo1 Birth Date: 1/1/1960 TPAIS Number: 123456789 Weight (kg): 55.00 Gender: Male Fm: Smt: Smt Onset: 3/1/1999 Yes																				
DA and Virus Load	<table border="1"> <tr> <td>DA</td> <td>30</td> </tr> <tr> <td>Ceftriaxone (mg)</td> <td>37</td> </tr> <tr> <td>Current Viral Load</td> <td>11</td> </tr> <tr> <td>Previous Viral Load</td> <td>17</td> </tr> </table> <p>Boundry and Prequalification Messages</p> <p>Plates & Reagents have been ordered and must be delivered by 12:00PM on 3/3/1999</p> <p>Specimen</p> <p>Specimen Type: V</p> <p>Poor Viral Suppression Δ: The patient's viral load count either did not decrease ≥ 5 log from the last point or is not below the viral load reduction goal. Unless lab error is at fault, consider changing therapy. More Info PQ1 p=QualA6, Commentary445</p> <p>Data Needed Soon - Caution</p> <ul style="list-style-type: none"> No Baseline Viral Load Value: Please specify which viral load value or values (an average of two point(s) you wish to be set as the baseline viral load value for this patient. BoundZY, Commentary411a <table border="1"> <tr> <td>Hemoglobin</td> <td>Specimen Date: V</td> </tr> <tr> <td>12</td> <td>3/1/1999</td> </tr> </table> <table border="1"> <tr> <td>Neutrophilic</td> <td>Specimen Date: G</td> </tr> <tr> <td>15</td> <td>3/1/1999</td> </tr> </table> <table border="1"> <tr> <td>Hepatic Function</td> <td>Specimen Date: A5</td> </tr> <tr> <td>49</td> <td>3/1/1999</td> </tr> </table>	DA	30	Ceftriaxone (mg)	37	Current Viral Load	11	Previous Viral Load	17	Hemoglobin	Specimen Date: V	12	3/1/1999	Neutrophilic	Specimen Date: G	15	3/1/1999	Hepatic Function	Specimen Date: A5	49	3/1/1999
DA	30																				
Ceftriaxone (mg)	37																				
Current Viral Load	11																				
Previous Viral Load	17																				
Hemoglobin	Specimen Date: V																				
12	3/1/1999																				
Neutrophilic	Specimen Date: G																				
15	3/1/1999																				
Hepatic Function	Specimen Date: A5																				
49	3/1/1999																				

FIG. 10D



TPMS 41

TPMS Patient # 1609 (70a)

Medical History Chart / Recent Evaluation

General		<input type="checkbox"/> Fatty	<input type="checkbox"/> Embo	<input type="checkbox"/> Compton	Date	Value
Patient Id	ARV naïve	Birth Date	TPMS Number	Weight (kg)	2/1/1999	73.00
Female				Solid/liquid	2/1/1999	Yes
CD4 and Viral Load		Specimen Date	Value	Specimen Date	Specimen Value	
CD4 Cells/ecl/mm ³	2/20/1999	350		2/20/1999	73	
CURRENT Viral Load	2/20/1999	31000	V. Urin	V. Urin	U/ml	
Previous Viral Load	12/31/1998	14000	V. Urin	V. Urin	U/ml	
HIV Genotype		Specimen Date	Value			
Phenotype						
Allergia/Hyper						
Intolerance						
Hemoglobin		Specimen Date	Value (Mg/dl)	Specimen Date	Value	
		2/1/1999	12.50			
Neutrophil		Specimen Date	Value (X10 ³ /mm ³)	Specimen Date	Value	
		2/1/1999	1350			
Hepatic Function		Specimen Date	Value (U/L)	Specimen Date	Value	
		2/1/1999	35			
AIDS Diagnosis		Specimen Date	Value	Specimen Date	Value	
Cytomegalovirus		Specimen Date	Value	Specimen Date	Value	
Epstein-Barr Virus		Specimen Date	Value	Specimen Date	Value	
Non-AIDS Drugs		Specimen Date	Value	Specimen Date	Value	
Tuberculosis		Specimen Date	Value	Specimen Date	Value	
Pozac Pulvules & Liquid, O... oral		Specimen Date	Value	Specimen Date	Value	
Beclim DS Tablets		Specimen Date	Value	Specimen Date	Value	
Relief Function		Specimen Date	Value	Specimen Date	Value	
Stomach Culture, E. Coli		Specimen Date	Value	Specimen Date	Value	
How To		Specimen Date	Value	Specimen Date	Value	

FIG. 11A

TPMS 41

How To

702

Medical History / Clinical History		HIVology Evaluation	
General		<input type="checkbox"/> Emphysema	<input checked="" type="checkbox"/> Comment/Pgroup
Patient ID	ARV naïve	Birth Date	1/5/1968
Physician		TRMIS Number	
		Weight (kg)	73.00
		Solid Dosage	27/1/1999
		Comments	Yes
CD4 and Viral Load		CD4 and Viral Load	
CD4 (cells/cubic mm)	500	CD4 (%)	27%
Cured Viral load	1	Cured Viral load (%)	27%
Previous Viral load	12	Comments	
HIV Genotype		HIV Genotype	
Genotype		Genotype	
Ale or Hyper		Ale or Hyper	
Tolerance		Tolerance	
Hemoglobin		Hemoglobin	
Specimen Date	2/1/1999	Specimen Date	12
Neutrophils		Neutrophils	
Specimen Date	2/1/1999	Specimen Date	13
Hepatic Function		Hepatic Function	
Specimen Date	ASAT	Specimen Date	35

• **Therapy Initiation:** Current treatment guidelines recommend initiation of antiRetroviral therapy for HIV-infected patients with HIV RNA (viral load) concentrations greater than 20,000 copies/ml (10,000 Eeq/ml bDNA) or CD4 counts less than 500 cellular (Ann. Int. Med., 1998). PreQualM, Commentary6

• **Combination Therapy Recommended:** Experts agree that the goal of antiRetroviral therapy should be to reduce the viral load to as low a level as possible for as long as possible. Initiation of therapy with a combination containing 2 nucleoside reverse transcriptase inhibitors (NRTI's) and a potent protease inhibitor have been shown to provide enhanced clinical benefit versus 2 drug combinations with regard to reduction in viral load and improved clinical outcomes. P=QualM, Commentary6

F16- 113

TPMS 41

How
To

WEB

W

S

E

P

C

D

F

G

TPMS Patient	
Medications	Chart
Retrovir 200mg q12h RTV only Evaluated	Initial Evaluation
	Initial Evaluation

Recommended Dosages

- Retrovir 300mg q12h (2 pills/day, \$9.56/day)
- Videl 200 mg q12h (4 pills/day, \$6.78/day)
- Norvir 600 mg q12h (12 pills/day, \$22.26/day)
- Resciptor 400 mg q8h (12 pills/day, \$7.39/day)

F16. 11D

- AZT: Interrupt Retrovirus if anemia and/or neutropenia develops. More Info 036 DosGenA, Commentary 36

CmtGenA, Commentary 13

- ddi: When treatment with other drugs known to cause pancreatic toxicity is required (for example, IV pentamidine), suspension of Videl should be considered.
- ddi: If patients develop symptoms of neuropathy, Videl therapy should be interrupted. DosGenB, Commentary 40 CmtGenA, Commentary 13
- ddi: Clinical signs suggestive of pancreatitis should prompt dose suspension of Videl and careful evaluation of the possibility of pancreatitis. Only after pancreatitis has been ruled out should dosing be resumed. DosGenB, Commentary 39

- DLV: Skin rash attributable to Resciptor may occur during first 21 days. More Info 054 CmtGenS, Commentary 54

- ddi: Videl should not be administered with a prescription antibiotic containing any form of tetracycline. CmtGenA, Commentary 5

- ddi: Plasma concentrations of some quinolone antibiotics are decreased when administered with antacids containing magnesium or aluminum. Therefore, doses of quinolone antibiotics should not be administered within 2 hours of taking Videl. CmtGenA, Commentary 16

- RTV: Monitor for decreased AUC of Norvir and associated adverse events when concomitant with use of drugs that increase CYP3A activity (including tobacco). More Info 026 CmtGenH, Commentary 26

TPMS

How
To

660

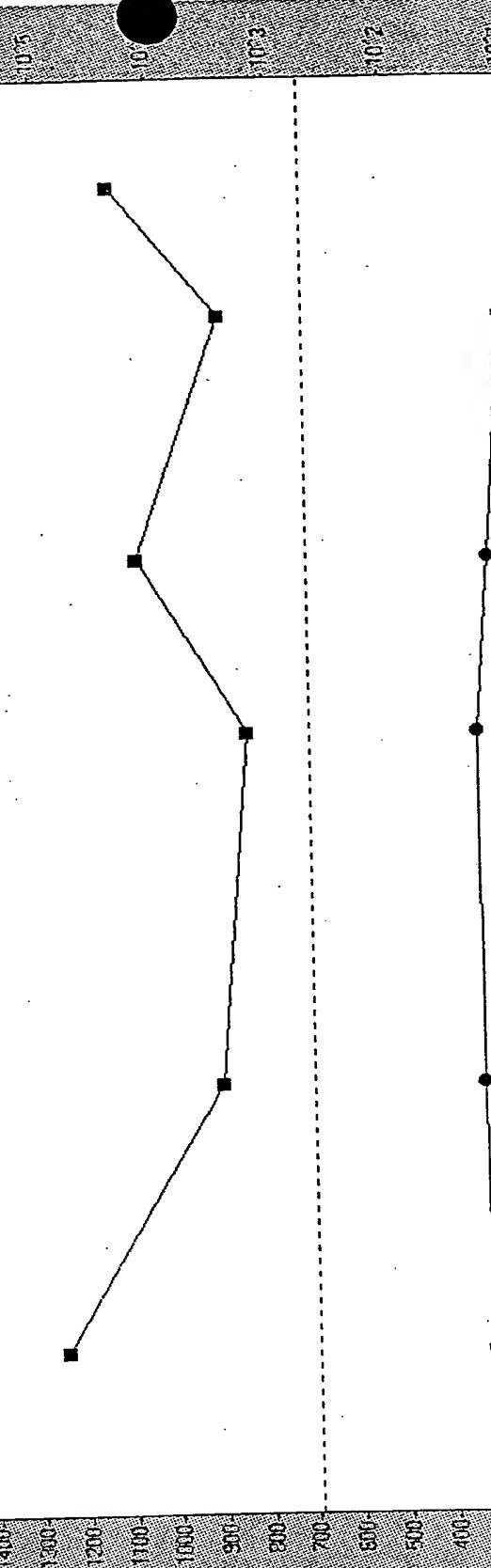
TPMS Patient

Written History Chart | Dose/Evaluation |

CD4 T Cells/cubic mm

1700
1500
1400
1300
1200
1100
1000
900
800
700
600
500
400
300
200
100
0

Fig. 12 A



A1 ↴

TPMS

Phenotypic Resistance to 3TC
from 3/15/1999 to present

dI

Nt

AZ

Mv

ddC

d4T

3TC

17/99

17/99

17/99

17/99

17/99

17/99

17/99

17/99

How
To

TPMS

TPMS Patient							
Patient History / Chart		Therapy Evaluation					
<input type="checkbox"/> Evaluate Clinical History <input checked="" type="checkbox"/> TC - ATV, NVP							
Initial Date: 10/6/2011		<input type="checkbox"/> Show 100% Predicted <input type="checkbox"/> Show 200% Predicted <input type="checkbox"/> Show 300% Predicted <input type="checkbox"/> Show 400% Predicted <input type="checkbox"/> Show 500% Predicted <input type="checkbox"/> Show 600% Predicted <input type="checkbox"/> Show 700% Predicted <input type="checkbox"/> Show 800% Predicted <input type="checkbox"/> Show 900% Predicted					
Severity: <input checked="" type="checkbox"/> Severe <input type="checkbox"/> Moderate <input type="checkbox"/> Minimal		<input type="checkbox"/> Antiviral Dose <input type="checkbox"/> Antiviral Adverse					
Therapy: <input checked="" type="checkbox"/> ddI, d4T, NFV <input checked="" type="checkbox"/> ddI, d4T, EFV <input type="checkbox"/> ddI, NFV, EFV <input type="checkbox"/> d4T, NFV, EFV <input type="checkbox"/> ddC, NFV, EFV <input checked="" type="checkbox"/> ddC, d4T, EFV		<input checked="" type="checkbox"/> AZT (Retrovir/zidovudine) <input type="checkbox"/> ddI (Videx/didanosine) <input type="checkbox"/> ddC (Hivid/zalcitabine) <input checked="" type="checkbox"/> 3TC (Epivir/lamivudine) <input checked="" type="checkbox"/> d4T (Zerit/stavudine) <input checked="" type="checkbox"/> ABC (Ziagen/abacavir)					
Genotype: See All <input checked="" type="checkbox"/> Print <input type="checkbox"/> Print Screen Evaluation		<input checked="" type="checkbox"/> Interactions <input checked="" type="checkbox"/> Mutations					
Therapy Rating: <input checked="" type="checkbox"/> N/A <input type="checkbox"/> Altered							
III THERAPY REJECTED !!! This therapy was rejected for the following reason(s) Additional information about the therapy is provided but this therapy is NOT advisable							
<p>• Viramune (nevirapine/NVP) Resistance Advisory: According to the last genotype data entered, the patient's virus currently has mutation(s) which is/are associated with resistance to Viramune. <input type="checkbox"/> Rejection54</p> <p>• Resistance Advisory: According to the last genotype data entered, the patient's virus currently has the following mutations; M184V [RT]. The genotype test displays evidence of the M184V/M184I mutation which is associated with resistance to 3TC. However, this mutant has increased sensitivity to the antiRetroviral activity of AZT and ADV so an AZT/3TC or AZT/ADV combination is still useable. Therefore combinations which contain AZT/3TC and AZT/ADV are shown as therapy options although these therapies have been ranked down +5 in favor of three drug combinations with no resistant mutants. <input type="checkbox"/> Rejection51</p> <p>• Epivir and Viramune Resistance Advisory: The patient's last phenotypic assay demonstrates phenotypic resistance to Epivir and Viramune, therefore, therapies containing Epivir and Viramune are not recommended at this time. <input type="checkbox"/> Rejection12</p>							
CAUTION YELLOW ALERT							
TPMS - 1							
• NVPΔ: Drug Interaction Alert: Patient is currently taking rifabutin and there is insufficient data to assess whether dose adjustments are necessary. These drugs							
✓✓✓ 3							

Medical History		Chart		Review Evaluation	
General					
Patient Id	Features1	Birth Date:	1/1/1960	TFNS#:	<input checked="" type="checkbox"/> Command Pending
Physician	patient	Gender:	Male	Weight (kg):	1/28/1999
				Date:	60.00
				Value:	
LTD and Vital Signs					
CMV	Specimen Date:	3/15/1999	Value:	<input checked="" type="checkbox"/> Freq:	<input checked="" type="checkbox"/> Comment Pending
(Cytomegalovirus)		240			
Current Viral Load		3/15/1999			
Previous Viral Load		21500			
		V.L Units:	C/ml		
		V.L Units:	U/ml		
		V.L Units:	U/ml		
HIV Genotype					
NVP	Specimen Date:	3/15/1999	Value:	<input checked="" type="checkbox"/> Y101[P], M461[P], Y541[P], Y82A[P], M411[R1], Y181	
(Nucleoside Reverse Transcriptase Inhibitor)					
Non-ARV Drugs					
<p>• NVPΔ: Drug Interaction Alert: Patient is currently taking rifabutin and there is insufficient data to assess whether dose adjustments are necessary. These drugs should only be used in combination if clearly indicated and with careful monitoring. CmdDP, Commentary 33</p>					
AIDS Diagnostic					
CD4	Specimen Date:	1/28/1999	Value:	<input checked="" type="checkbox"/> Date:	<input checked="" type="checkbox"/> Comment Pending
(Cellular Immunity)		75			
Neurology					
Paroxysmal	Specimen Date:	1/28/1999	Value:	<input checked="" type="checkbox"/> Date:	<input checked="" type="checkbox"/> Value:
(Paroxysmal Seizures)		15.00		1/28/1999	No
Hematology					
Neutrophil	Specimen Date:	1/28/1999	Value:	<input checked="" type="checkbox"/> Date:	<input checked="" type="checkbox"/> Value:
(Neutrophils)		1500		1/28/1999	No
Renal Function					
Urea	Specimen Date:	1/28/1999	Value:	<input checked="" type="checkbox"/> Date:	<input checked="" type="checkbox"/> Value:
(Plasma Urea Nitrogen)		150		1/28/1999	No
Hepatic Function					
Transaminase	Specimen Date:	1/28/1999	Value:	<input checked="" type="checkbox"/> Date:	<input checked="" type="checkbox"/> Value:
(Plasma Transaminase)		25		1/28/1999	No

Fig. 12 C

How to... WEB